A Alkaline Peptone Water, cont

Limitations of the Procedure

This prepared tube medium is intended to be used as an enrichment medium. A pure culture is recommended for biochemical tests and other identification procedures.

References

- Gilligan, Janda, Karmali and Miller. 1992. Carairech 12A, Laboratory diagnosis of bacterial disreters. Goord. ed., Nobte. American Secrety for Microbiology, Washington, D.C. Foebes, Sahm and Weissfeld. 1998. Balley & Scott's diagnostic microbiology, 10th ed. Mosthy, Inc. St. Louis, Mo.
- LOUIS, MO.
 Grassuck. 1992. In Isenberg (ed.), Clinical nucrobiology procedures handbook, vol. 1. American Society for Microbiology, Washington, D.C.

Availability

BBL™ Alkaline Peptone Water

Cat. No. 297814 Prepared Tubes, 8 ml. (D Tubes) - Pkg. of 10

Amies Transport Media

(See Transport Media)

Amino Acid Assay Media Lysine Assay Medium • Methionine Assay Medium **Cystine Assay Medium**

Intended Use

Lysine Assay Medium is used for determining lysine concentration by the microbiological assay technique.

Methionine Assay Medium is used for determining methionine concentration by the microbiological assay technique.

Cystine Assay Medium is used for determining L-cystine concentration by the microbiological assay technique.

Summary and Explanation

Amino acid assay media are prepared for use in the microbiological assay of amino acids. Three types of media are used for this purpose:

- 1. Maintenance Media: For carrying the stock culture to preserve the viability and sensitivity of the test organism for its intended purpose:
- 2. Inoculum Media: To condition the test culture for immedi-
- 3. Assay Media: To permit quantitation of the amino acid under test. They contain all the factors necessary for optimal growth of the test organism except the single essential amino acid to be determined.

Amino Acid Assay Media are prepared according to the formulations of Steel et al.1 They are used in the microbiological assay of amino acids using Pediococcus acidilactici ATCC" 8042 as the test organism.

Principles of the Procedure

Lysine Assay Medium, Methionine Assay Medium and Cystine Assay Medium contain all the factors essential for the growth of Pediococcus acidilactici ATCC 8042, except the amino acid under assay. The addition of the amino acid in specified increasing concentrations gives a growth response by the test organism.

Formulae¹

Difco™ Lysine Assay Medium, Methionine Assay Medium or Cystine Assay Medium

Approximate Formula* Per Liter

All amino acid assay media contain the following formula.

Sodium Acetate40.0 Monopotassium Phosphate1.2 Ferrous Sulfate Manganese Sulfate40.0 mg Sodium Chloride20,0 mg Guanine Hydrochloride20.0 mg Yanthine Thiamine Hydrochloride 1.0 mg Pyrodoxine Hydrochloride2.0 mg Riboflavin1.0 mg p-Aminobenzoic Acid200.0 μg L-Histidine Hydrochloride124.0 mg DL-Phenylalanine 0.2 L-Arginine Hydrochloride484.0 mg

User Quality Control

Identity Specifications

Difco" Lysine Assay Medium, Methionine Assay Medium or Cystine Assay Medium

Dehydrated Appearance: White to off-white, homogeneous, may have a tendency to clump.

Solution: 5.25% (single strength) solution, soluble in purified water upon boiling. Solution is light to medium amber, clear, may have a slight precipitate;

Prepared Appearance: Single strength—Light to medium amber, clear, may have a slight precipitate

Reaction of 5.25% Solution at 25°C: pH 6.7 ± 0.2

Cultural Response

Difco" Lysine Assay Medium, Methionine Assay Medium or Cystine Assay Medium

Prepare the medium per label directions. These media support the growth of Pediococcus additactic/ATCC* 8042 when prepared in single strength and supplemented with the appropriate amino acid, upine Assay Medium should produce a standard curve when tested with L-typine at 0.0 to 300 upper 10 mil. Methioline Assay Medium should produce a standard curve when tested with L-typine Assay Medium should produce a standard curve when tested with L-typine at 0 to 50 upper 10 mil. Incubate tubes with caps lossened at 35-37°C for 16-20 hours? Read the percent transmittance at 660 mil.

Preparation of inoculum dilution, amino acid stock and working solution.

PREPARATION OF INCCULUM DILUTION PREPARATION OF (CELL SUSPENSION + AMBIO ACID STOCK SOL ASSAY MEDIUM TEST CULTURE STERILE 0.35% NACI) (AMINO ACID + PURPHE	UTION (STOCK SOLUTION + SOLUTION CONCENTRATION
. Medium ATCC** 8042 Methionine Assay Pediococcus acidifactio 1 mL + 19 mL DL-methionine 1.2 g + 1. Medium ATCC** 8042	2, 2.5, 3, 4, 5 150, 180, 240, 300 ,000 mt 1 mt + 99 mt 0, 0.5, 1, 1.5, 0.0, 6, 12, 18, 24, 2, 2.5, 3, 4, 5 30, 36, 48, 60

In addition to the ingredients listed on the previous page, the media contain per liter*:

Lysine Assay Medium	
L-Cystine	0.1 g
DL-Methionine	0.2 g
Methionine Assay Medium	
L-Cystine	0.1 g
L-Lysine Hydrochloride	0.5 g
Cystine Assay Medium	
DL-Methionine	0.2 a
L-Lysine Hydrochloride	0.5 g
*Adjusted and/or supplemented as required to meet performance criteria.	

Precautions

Great care must be taken to avoid contamination of media or plassware in microbiological assay procedures. Extremely small amounts of foreign material may be sufficient to give erroneous results. Scrupulously clean glassware free from detergents and other chemicals must be used. Glassware must be heated to 250°C for at least 1 hour to burn off any organic residues that might be present. Take precautions to keep sterilization and cooling conditions uniform throughout the assay.

Directions for Preparation from Dehydrated Product

completely dissolve the powder.

- Suspend 10.5 g of the powder in 100 mL of purified water.
 Heat with frequent agitation and boil for 2-3 minutes to
- Dispense in 5 mL amounts into tubes, evenly dispersing the precipitate.

- 4. Add standard or test samples.
- Adjust tube volume to 10 mL with purified water.
- 6. Autoclave at 121°C for 10 minutes.

Procedure

Stock Culture and Inoculum

Stock cultures of Pediococcus acidilactici ATCC 8042 are prepared by stab inoculation into tubes of Lactobacilli Agar AOAC or Micro Assay Culture Agar. Incubate cultures at 35-37°C for 24 hours. Store stock cultures at 2-8°C. Make transfers at monthly intervals in triolicate.

The inoculum for assay is prepared by subculturing the test organism into 10 mL Latobacilli Broth AOAC or Micro Inoculum Broth. Incubate at 35-37°C for 16-24 hours. After incubation, centrifuge the cells under aseptic conditions and decant the liquid supernatun. Wash the cells 3 times with 10 mL sterile 0.85% NaCl solution. After the third wash, resuspend the cells in 10 mL sterile 0.85% NaCl solution. Dilute the 10 mL cell suspension with the appropriate amount of sterile 0.85% NaCl solution. (See the table under User Quality Control, Cultural Response.) One drop of the diluted inoculum suspension is used to inoculate each of the assay tubes

Amino Acid Solution

Prepare stock solutions of each amino acid as described in Table 1. If the DL form is used, twice the concentration of the amino acid is required. Prepare the stock solutions fresh daily.